UNITED STATES DEPARTMENT OF THE INTERIOR GEOLOGICAL SURVEY

Analytical results and sample locality map
of heavy-mineral-concentrate and rock samples
from the Organ Mountains Wilderness Study Area (NM-030-074),
Dona Ana County, New Mexico

Ву

Tracy A. Delaney¹, Gordon W. Day², Robert L. Turner¹, and Janet L. Jones¹

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This report is preliminary and has not been reviewed for conformity with U.S. Geological Survey editorial standards and stratigraphic nomenclature. Any use of trade names is for descriptive purposes only and does not imply endorsement by the USGS.

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STUDIES RELATED TO WILDERNESS

Bureau of Land Management Wilderness Study Areas

The Federal Land Policy and Management Act (Public Law 94-579, October 21, 1976) requires the U.S. Geological Survey and the U.S. Bureau of Mines to conduct mineral surveys on certain areas to determine their mineral values, if any. Results must be made available to the public and be submitted to the President and the Congress. This report presents the results of a geochemical survey of the Organ Mountains Wilderness Study Area (NM-030-074), Dona Ana County, New Mexico.

INTRODUCTION

In the fall of 1984, the U.S. Geological Survey conducted a reconnaissance geochemical survey of the Organ Mountains Wilderness Study Area, Dona Ana County, New Mexico.

The Organ Mountains Wilderness Study Area (WSA) comprises about 11 mi^2 (28.5 km²) (7,283 acres) in the eastern Dona Ana County, New Mexico. The study area is located about 15 miles east-northeast of Las Cruces, New Mexico (fig. 1).

The WSA has a semi-arid climate and is drained by intermittent streams. Ponderosa pine is the dominant vegetation on the upper part of the mountains with pinyon pine and juniper at lower elevations. The lower mountain slopes contain mountain mahogany, oak, mesquite, and cresote shrubs.

The topographic relief in the study area is about 3000 ft (914 m), with a maximum elevation of 8010 ft (2441 m). Access to the study area is obtained by U.S. highway 70 on the north side, a graveled road on the west side, and a paved road on the east side.

The Organ Mountains WSA is situated within the Basin and Range Physiographic Province. The major portion of the study area is a quartz monzonite batholith of Tertiary age which was intruded into Paleozoic sedimentary rocks (limestone, dolomites, and shales).

METHODS OF STUDY

Sample Media

Heavy-mineral-concentrate samples provide information about the chemistry of certain minerals in rock material eroded from the drainage basin upstream from each sample site. The selective concentration of minerals, many of which may be ore related, permits determination of some elements that are not easily detected in stream-sediment samples.

Analyses of unaltered or unmineralized rock samples provide background geochemical data for individual rock units. On the other hand, analyses of altered or mineralized rocks, where present, may provide useful geochemical information about the major- and trace-element assemblages associated with a mineralizing system.

Sample Collection

Heavy-mineral-concentrate samples were collected at 17 sites (plate 1). Twelve rock samples were collected at 8 sites. Sampling density was about one sample site per 0.65 mi² for the heavy-mineral concentrates and about one

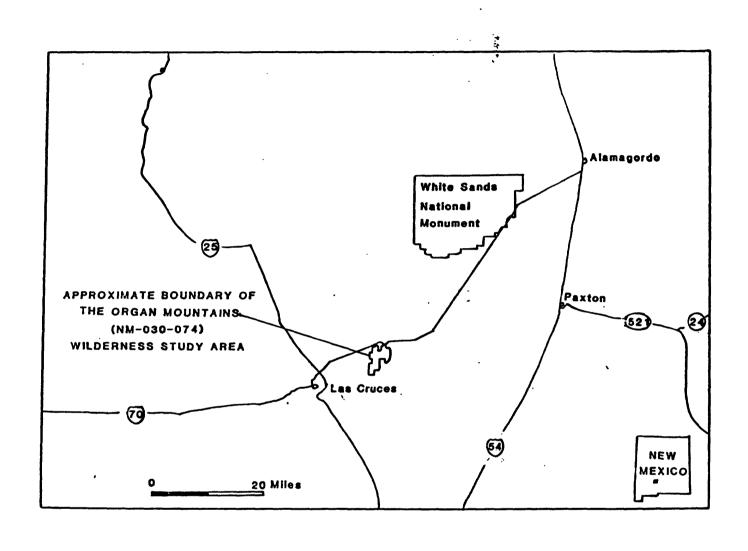


Figure 1. Location map of the Organ Mountains Wilderness Study Area (NM-030-074), Dona Ana County, New Mexico.

sample per 1.38 mi^2 for the rocks. The area of the drainage basins sampled ranged from .25 mi^2 to 4 mi^2 .

Heavy-mineral-concentrate samples

Heavy-mineral-concentrate samples were collected from active alluvium. Each bulk sample was screened with a 2.0-mm (10-mesh) screen to remove the coarse material. The less than 2.0-mm fraction was panned until most of the quartz, feldspar, organic material, and clay-sized material were removed.

Rock samples

Rock samples were collected from various types of occurrences in the vicinity of the plotted site location. Descriptions of rock samples are in table 5.

Sample Preparation

After the samples were air dried, bromoform (specific gravity 2.8) was used to remove the remaining quartz and feldspar from the heavy-mineral-concentrate samples that had been panned in the field. The resultant heavy-mineral sample was separated into three fractions using a large electromagnet (in this case a modified Frantz Isodynamic Separator). The most magnetic material, primarily magnetite, and the second fraction, largely ferromagnesian silicates and iron oxides, were saved for archival storage. The third fraction (the least magnetic material which may include the nonmagnetic ore minerals, zircon, sphene, etc.) was split using a Jones splitter. One split was hand ground for spectrographic analysis; the other split was saved for mineralogical analysis. These magnetic separates are the same separates that would be produced by using a Frantz Isodynamic Separator set at a slope of 15° and a tilt of 10° with a current of 0.2 ampere to remove the magnetite and ilmenite, and a current of 0.6 ampere to split the remainder of the sample into paramagnetic and nonmagnetic fractions.

Rock samples were crushed and then pulverized to minus $0.15\ \mathrm{mm}$ with ceramic plates.

Sample Analysis

Spectrographic method

The heavy-mineral-concentrate and rock samples were analyzed for 31 elements using semiquantitative, direct-current arc emission spectrographic methods. The analyses for heavy-mineral-concentrate samples were performed by analysts in the Branch of Exploration Geochemistry using the method of Grimes and Marranzino (1968); analyses for rock samples were performed by analysts in the Branch of Analytical Chemistry using a modified method of Myers and others (1961) by Crock and others (1987). The elements analyzed and their lower limits of determination are listed in table 1. For arsenic (As), gold (Au), cadmium (Cd), lanthanum (La), and thorium (Th), the lower limits of determination of the two analytical methods varies. The values in the parentheses are the limits of determination for Myers and others (1961). Spectrographic results were obtained by visual comparison of spectra derived from the sample against spectra obtained from standards made from pure oxides and carbonates. Standard concentrations are geometrically spaced over any

given order of magnitude of concentration as follows: 100, 50, 20, 10, and so forth. Samples whose concentrations are estimated to fall between those values are assigned values of 70, 30, 15, and so forth. The precision of the analytical method is approximately plus or minus one reporting interval at the 83 percent confidence level and plus or minus two reporting intervals at the 96 percent confidence level (Motooka and Grimes, 1976). Values determined for the major elements, iron, magnesium, calcium, and titanium, are given in weight percent; all others are given in parts per million (micrograms/gram). Analytical data for samples from the Organ Mountains WSA are listed in tables 3 and 4.

Chemical methods

Other analytical methods used on rock samples from the Organ Mountains Wilderness Study Area are summarized in table 2. The analytical method used for determining As, Bi, Cd, Sb, and Zn is a modification and adaptation by Crock and others (1987) for the inductively coupled plasma method (ICP) based on the method of O'Leary and Viets (1986).

Analytical results for rock samples are listed in tables 3, and 4, respectively.

ROCK ANALYSIS STORAGE SYSTEM

Upon completion of all analytical work, the analytical results were entered into a computer-based file called Rock Analysis Storage System (RASS). This data base contains both descriptive geological information and analytical data. Any or all of this information may be retrieved and converted to a binary form (STATPAC) for computerized statistical analysis or publication (VanTrump and Miesch, 1977).

DESCRIPTION OF DATA TABLES

Tables 3 and 4 list the results of analyses for the samples of heavy-mineral concentrate and rock, respectively. For the two tables, the data are arranged so that column 1 contains the USGS-assigned sample numbers. These numbers correspond to the numbers shown on the site location map (plate 1). Columns in which the element headings show the letter "s" below the element symbol are emission spectrographic analyses; "icp" indicates inductively coupled plasma-atomic emission spectroscopy. A letter "N" in the tables indicates that a given element was looked for but not detected at the lower limit of determination shown for that element in table 1. If an element was observed but was below the lowest reporting value, a "less than" symbol (<) was entered in the tables in front of the lower limit of determination. If an element was observed but was above the highest reporting value, a "greater than" symbol (>) was entered in the tables in front of the upper limit of determination. If an element was not looked for in a sample, two dashes (--) are entered in tables 3 and 4 in place of an analytical value. Because of the formatting used in the computer program that produced tables 3 and 4, some of the elements listed in these tables (Fe, Mg, Ca, Ti, Ag, and Be) carry one or more nonsignificant digits to the right of the significant digits. The analysts did not determine these elements to the accuracy suggested by the extra zeros.

ACKNOWLEDGMENTS

A number of our colleagues also participated in sample collection, preparation and analyses of these samples: collection, Allen Phillips and Rick Graff; preparation, Tom Peacock; and analyses, Mollie Malcolm and Paul Briggs.

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TABLE 1.--Limits of determination for the spectrographic analysis of rocks based on a 10-mg sample

[The values shown are the lower limits of determination assigned by the Grimes and Marranzino method, except for those values in parentheses, which are the lower values assigned by the Myers and others method. The spectrographic limits of determination for heavy-mineral-concentrate samples are based on a 5-mg sample, and are therefore two reporting intervals higher than the limits given for rocks.]

		Percent	
Iron (Fe) Magnesium (Mg) Calcium (Ca) Titanium (Ti)	0.05 .02 .05 .002		20 10 20 1
	Part	s per million	
Manganese (Mn) Silver (Ag) Arsenic (As) Gold (Au) Boron (B) Barium (Ba) Beryllium (Be) Bismuth (Bi) Cadmium (Cd) Cobalt (Co) Chromium (Cr) Copper (Cu) Lanthanum (La) Molybdenum (Mo) Niobium (Nb) Nickel (Ni) Lead (Pb) Antimony (Sb) Scandium (Sc) Tin (Sn) Strontium (V) Tungsten (W) Yttrium (Y) Zinc (Zn) Zirconium (Zr)	10 0.5 200 10 10 20 1 10 20 5 10 5 20 5 10 100 5 10 100 100 100 100	(700) (15) (30)	5,000 5,000 10,000 500 2,000 5,000 1,000 500 2,000 5,000 2,000 1,000 2,000 2,000 10,000 10,000 10,000 10,000 10,000 10,000 11,000 1,000 11,000 11,000

TABLE 2.--Chemical methods used

[ICP = inductively coupled plasma spectroscopy]

Element or constituent determined	Sample type	Method	Determination limit (micrograms/ gram or ppm)	Reference
Arsenic (As)	rock	ICP	5	Crock and others,
Antimony (Sb)	rock	ICP	2	1987.
Zinc (Zn)	rock	ICP	2	
Bismuth (Bi)	rock	ICP	2	
Cadmium (Cd)	rock	ICP	0.1	

	TABLE	т СО	3 RESULTS OF ANALYSES OF	ILTS 0	F A	IALYSES	Ī	EAVY-MINERAL-CONCENTRATE SAMPLES FROM THE ORGAN MOUNTAINS BLM WILDERNESS STUDY ABEAL DONA AND COUNTY, NEW MEYICO	AL-CONCENTRATE SAMPLES FROM	S FROM THE OF	GAN MOUNTA	NS BLM WIL	DERNESS STU	,
IN, not detected; (, detected but below th	90	80 t e	÷	etect	D •	ut be	=		tion shown;	determination shown;), determined to be greater than the value shown.]	ed to be gre	ater than	the value sl	[· uwor
Sample	۲.	Latitude	•	Long	Long! tude	•	Fe-pot.	Mg-pot.	Ca-pot.	Ti-pot.	Maga-nM s	Ag-ppm 8	As-ppm	Au-ppm
OM0 0 1H	35	22 5	53	106	32 5	58	-	1.00	e.	-	100	z	Z	z
OM0 0 2 H	32	23	_		32 5	-	. 2	. 10	1.0	۲.	0.00	z	z	z
HE 0 0WO	32	23	9	106		33	,2	.03	.	۲.	00	z	z	z
OM0 0 4 H	32	23	57		32 2	9	.2	. 0.5	2.0	٠. د	00-	z	z	z
OM0 0 3H	32	24 2	9	106	33	4	. 2	.03	o. n	٠.	130	z	z	z
OM0 0 6H	32	24 2	29	106	66	m	8.	0.	20.0	1.0	300	z	z	z
OM0 0 7 H	32	25 3	38	106	35 2	*		50	2,0	٠.	9	'n	z	z
OM0 0 8H	32		=	106	36	52	ю.	.30	2.0	٠.	200	z	z	z
H600W0	32	21 4	80	90+	•	47		.30	0.0	.2	130	z	z	z
OM0 1 0 H	32	22	'n	106	36	en En		. 10	1.0	1.0	100	z	z	z
OM0 1 1 H	32	22	6	106	36	6		.20	1.0	'n	200	z	z	z
OM0 12H	32	23	12	106	36	67	0.1	.20	۲.	0.5	200	z	z	z
OM0 13H	32	23 3	38	406	37	0	'n	0.	2.0	2.0	200	z	z	z
OM0 14H	32	23 5	55	106	37	2	٠.	. 50	7.0	2.5	300	~	z	z
OM0 15H	32	24 4	9	106	36 4	• •	۲,		o.	. 7	100	z	z	z
OM0 1 6H	32	24	6 4		96	Ξ	7.	.03	0.0	2.0	150	z	z	z
OM0 17H	32	24	52	106	36	4	e,	. 30	o. v	r	200	z	z	z

	TABLE 3	RESULTS OF ANALYSES OF	ANALYSES OF		HEAVY-MINERAL-CONCENTRATE SAMPLES FROM THE ORGAN MOUNTAINS AREA, DONA ANA COUNTY, NEW MEXICOContinued	TRATE SAMPLY, NEW MEXI	ES FROM T COConti	HE ORBAN I	MOUNTAINS		BLM WILDERNESS STUDY	
Samole	mdd- i N	Pb-ppm	Edd-q8	Edd-og	Edd-uS	Sr-ppm	Edd-∧	E 4 - >	Mqq-Y	Zn-ppm	Zr-ppm	Th-ppm
	•	•	•	•	•	•	•	•	•	•	•	•
OM0 0 1 H	z	100	z	10	z	200	(20	< 100	00+	z	12,000	Z
OM0 0 2H	z	00+	z	0	z	200	20	z	130	z)2,000	300
OMO OH	z	z	z	9	z	z	(20	z	150	z	72,000	z
OM004H	z	20	z	9-	z	z	(20	300	150	z	72,000	300
OMO 0 SH	z	100	z	10	z	200	20	300	200	z	12,000	900
OM0 0 6H	z	z	z	20	z	z	20	100	200	z)2,000	z
OM0 07H	z	200	z	-	z	200	20	100	150	z	72,000	z
OM0 0 BH	z	20	z	20	z	300	(20	z	200	z	72,000	z
H600W0	z	200	z	2	z	z	(20	z	150	z	72,000	z
OM0 1 0 H	z	z	z	20	z	z	(20	Z	200	z)2,000	z
OM0 11H	z	100	z	5	z	z	(20	z	20	z	12,000	z
OM0 12H	z	0,2	z	20	z	z	20	z	300	z	72,000	Z
OM0 13H	z	20	z	30	z	z	20	z	700	Z	72,000	Z
OM0 14H	z	1,000	z	30	z	z	30	z	300	3,000)2,000	Z
OM6 15H	z	200	z	90	z	z	(20	z	300	z)2,000	z
OM0 16H	z	300	z	30	z	z	20	z	700	1,000)2,000	z
OM0 17H	z	200	z	20	z	z	20	z	300	1,000	>2,000	z

TABLE 4 -- RESULTS OF ANALYSES OF ROCK SAMPLES COLLECTED FROM THE ORGAN MOUNTAIN BLM MILDERNESS STUDY AREA, DONA ANA

7	80 	1,500	300	200	300	70	70	20	20	70	420	6 6 9
ue shown	8 60 8 8	410 410	<10	<10	<10	¢10	~10	<10	<10	<10	<10	~10
an the va	AC-104	<15 <15	<15	<15	<15	<15	<15	<15	<15	<15	<15	<15
rester th	A 60 60 60 60 60 60 60 60 60 60 60 60 60	<700 <700	<700	<700	< 100	<700	<700	700	700	1,500	<700	< 100
d to be g	Ag-pp.	A A N N	5.	1.0	150.0	20.0	2.0	300.0	150.0	100.0	30.0	4. 5
determine	E G G G	150	0 M	10	100	>5,000	70	2,000	100	150	2,000	1,500
MEXICO shown; >, determined to be greater than the value shown.]	Ti-pct.	.500	100	.150	• 500					500.		-002
IN, not detected; <, detected but below the limit of determination a	Ca-pct.	.70	•15	•10	.10	1.50	08.	2.00	3.00	•30	7.00	15.00
limit of c	M9-pct.	2.00	50.	.15	.30	.70	<.02	.03	-02	<.02	.15	7.00
below the	Fe Boct.	3.00	30	7.00	10.00	10.00	.15	• 10	• 50	3.00	1.00	2.00
detected but	Longitude	106 32 57 106 32 57	32	33	32	106 36 0	35	35		32	106 35 51	32
detected; <,	Latitude	32 23 11 32 23 11	23	24	52	32 24 8	23	23		23	32 23 55	32 22 2
IN, not	Semon	OM002R1	OM004R	OM005R1	OM007R1	OM018R	OM019R	OM020R1	OM020R2	OM020R3	OM020R4	UM021R

TABLE 4 -- RESULTS OF ANALYSES OF ROCK SAMPLES COLLECTED FROM THE ORGAN MOUNTAIN BLM WILDERNESS STUDY AREA, DONA ANA COUTY, NEW MEXICO--Continued

SC-ppm	v û í	\$ \$ \$	A A A A A A A A A A A	ۍ ئ
86-pp#	0000	300	4100 1,000 4100 2,000	1,500
Pag- 94	410 610	700	7,000 30 15,000 20,000	7,000
- X - C C C C C C C C C C C C C C C C C C C	Ĉ.) - \$	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	î.
# dd - q N	00 00 00 00 00 00 00 00 00 00 00 00 00	420 420 420	00000 750 750 750 750	420 420
.M 	r v v	150	20 200 7 7	100
	V V V V V V V V V V V V V V V V V V V	200	00000 MMMMM V V V V	<30 <30
E C C	300 70	700	200 3,000 1,000 3,000	>20,000 300
	0 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	200		4 10 0 10 0 10 0 10 0 10 0 10 0 10 0 10
Co 1 pp	∾ လ ពុ	, ու Ն	& & & & & &	\$ \$
J	< 30 < 30 < 30	200 430 430 430 430 430 430 430 430 430 4	4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	100
84 - 100 m	000	700		0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
8 9 8 8	1.5	N - 0	00000	^!. ^!.0
Semole	OM002R1	0M005R1	0M018R 0M019R 0M020R1 0M020R2	OM020R4 OM021R

TABLE 4 -- RESULTS OF ANALYSES OF ROCK SAMPLES COLLECTED FROM THE ORGAN MOUNTAIN BLM WILDERNESS STUDY AREA, DONA ANA COUNTY, NEW MEXICO--Continued

Zn-ppm fcp	4W 40 4M 60	13,000 29 10,000 2,400	34,000
Sb-ppm icp	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	21 660 62 1,500	1,400
Cd-ppm icp	# W F W	61.0 86.0 1.0 7.7	73.0
81-ppm 1cp	<22 <22 <2 12 740	4 % % % %	m Q
As-ppm 1cp	% C C C C C C C C C C C C C C C C C C C	330 455 760 1,400	790
Th-ppm	00000 0000 0000 0000 0000 0000 0000	00000000000000000000000000000000000000	<200 <200
Zr-ppm	300 300 150 100	V V V V V V V V V V V V V V V V V V V	4 to 0
Zn-ppm	00000 00000 00000	>10,000 >200 >10,000 3,000	>10,000 700
F00*	A 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	00000	410 410
X 00 % E	A A A A A A A A A A A A A A A A A A A	4 4 5 5 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	<50
		00000 10000	<10 15
ed-18	500 100 150 100	* * * * * * * * * * * * * * * * * * *	4100 4100
Edd-rS 8	, , , , , , , , , , , , , , , , , , ,	7 V V V V V V V V V V V V V V V V V V V	15
Sample	OMO 0 2 R 1 OMO 0 2 R 2 OMO 0 4 R OMO 0 5 R 1	OM018R OM019R OM020R1 OM020R2	0M020R4 0M021R

Table 5. Description of rock samples

OM018R	mineralized rock
19R	siliceous
20R1	quartz vein
20R2	guartz vein
20R3	guartz vein
20R4	guartz vein
21R	mineralized rock from dump
OM002R1	quartzite
2R2	iron altered quartz monzonite
4R	altered quartz monzonite
5R1	iron-stained quartz monzonite
7R1	altered quartz monzonite